

Art Unit: ***

CLMPTO

Diane Williams

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Claims 1 – 7 have been canceled

8. A DNA sequence which codes for a protein having an enzymatic activity of orotate phosphoribosyl-transferase (pyrF activity) which comprises

a DNA sequence selected from the group consisting of the DNA sequence SEQ ID NO: 1 in a region from position 1133 up to and including position 1877,

the DNA sequence SEQ ID NO: 2 in a region from position 1 up to and including position 684,

a DNA sequence having a sequence homology of more than 70% with the said region of the DNA sequence SEQ ID NO: 1, and

a DNA sequence having a sequence homology of more than 70% with the said region of the DNA sequence SEQ ID NO: 2.

9. A protein having pyrF activity, which comprises an amino acid sequence selected from the group consisting of

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the amino acid sequence SEQ ID NO: 3; and
an amino acid sequence having a sequence homology of
more than 70% with the amino acid sequence SEQ ID NO: 3.

10. An expression vector which comprises a DNA sequence as
claimed in claim 8.

11. A microorganism which comprises an expression vector as
claimed in claim 10.

12. A process for producing fungal strains which are
capable of efficient expression and secretion of proteins,
comprising

transforming a fungal strain with an auxotrophic gene
defect as host strain in a transformation mixture, using with an
expression vector which has a gene for complementation of the
auxotrophic gene defect in the host strain;

selecting clones transformed with the expression vector
from the transformation mixture by selection for complementation
of the auxotrophic gene defect;

Art Unit: ***

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controlling expression of the gene for complementation of the auxotrophic gene defect in the host strain by a genetic regulatory element which is active in the host strain; and

employing as host strain a uridine-auxotrophic fungus selected from the group of genera consisting of *Trametes*, *Coriolus* and *Polyporus* with a gene defect in the *pyrF* gene.

13. An expression system comprising

a host strain selected from the group of genera consisting of *Trametes*, *Coriolus* and *Polyporus* having a genetic defect in metabolism, on the basis of which the metabolite uridine which is essential for growth is no longer synthesized, and the host strain is no longer able to grow on minimal media without addition of this metabolite; and

an expression vector comprising a selection marker gene which complements the auxotrophic gene defect of the host strain, wherein the host strain has a genetic defect in metabolism a defect in the *pyrF* gene, and the selection marker gene is the *pyrF* gene from a fungus of the class Basidiomycetes.

Art Unit: ***

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14. A process for producing a protein, which comprises
employing an expression system as claimed in claim 13
comprising a gene encoding the protein in a manner known in a
culture for protein production; and
obtaining the protein from the culture.

15. A process for producing a protein, which comprises
cultivating in a culture a microorganism as claimed in
claim 11, comprising a gene encoding the protein; and
obtaining the protein from the culture.

16. A process for producing a protein, which comprises
cultivating in a culture a fungal strain produced by a
process as claimed in claim 12, comprising a gene encoding the
protein; and
obtaining the protein from the culture.